

6/30/99

Washington State Medical Test Site Rules
PRE-INSPECTION SELF-ASSESSMENT CHECKLIST

TESTING IN DERMATOLOGY PRACTICE
MICROSCOPIC TISSUE EXAMS, KOH PREPS, DERMATOPHYTE CULTURES

I. MICROSCOPIC TISSUE EXAMS

PERSONNEL - High complexity testing

- ☐ M.D. and board certified in dermatology, dermatopathology or anatomic pathology
- ☐ Documentation of continuing education activities (i.e., case reviews at professional meetings; documentation of reviews by consultants; participation in proficiency testing; etc.)

RECORDKEEPING

- ☐ Charted order has all pertinent patient and specimen information
- ☐ Requisition form sent to tissue processing lab has: patient name or identifier; name of person ordering the test; date of collection; age; sex; other pertinent patient/specimen information
- ☐ Accession log or other system to track tissues sent out for processing:
 - ☐ Date specimen collected
 - ☐ Date processed/stained
 - ☐ Date slides reviewed
 - ☐ Date reported/charted
- ☐ Specimen labeling is:
 - ☐ Adequate on tissue containers, blocks, slides
 - ☐ Adequate to identify specimen for the required retention limit
- ☐ A system is in place to assure slides come back from the processing lab
- ☐ A system is in place for charting/recording results of microscopic exam
- ☐ A system is in place for notification of patients of abnormal results
- ☐ Reports are readily accessible
- ☐ Specimen limitations are noted where applicable
- ☐ A system is in place for corrected reports
- ☐ There is documentation of consult reviews

RETENTION

- ☐ Specimen blocks are:
 - ☐ Retained for 2 years
 - ☐ Stored under proper conditions
- ☐ Slides are retained for 10 years
- ☐ Reports are retained for 10 years
- ☐ Accession logs, requisitions are retained for 2 years

QUALITY CONTROL/QUALITY ASSURANCE

- ☐ Written procedures/policies for specimen collection, handling, preservation, labeling, referral for processing, retrieval of slides, review, reporting
- ☐ Review of quality control slides for special stains
- ☐ Documentation of consults, proficiency testing, case study reviews, professional meetings, other continuing education activities

SAFETY

- ☐ Policies for handling of specimens, infectious waste
- ☐ Policies for handling, storage, disposal of hazardous chemicals

DERMATOLOGY PRACTICE TESTING CHECKLIST

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II. KOH PREPS, DERMATOPHYTE CULTURES (Growth/No Growth Only)

PERSONNEL - Moderate complexity testing

- ___ Documentation of training and experience for testing performed
- ___ Written job description
- ___ Documentation of assessment of competency
- ___ Records of continuing education
- ___ Participation in proficiency testing or biannual verification of accuracy activities (2 samples, 2 times per year)

QUALITY CONTROL

- ___ Procedures written for KOH preps including: specimen collection and handling; preparation of reagents; preparation and examination of slides; interpretation of results; reporting protocol.
- ___ Procedures written for dermatophyte cultures, including: specimen collection and handling; inoculation of media; incubation requirements; review of growth and interpretation of results; reporting protocol. (Manufacturer's product inserts may be used)
- ___ Reagents and media are properly labeled, stored and within expiration date
- ___ For commercially prepared media, manufacturer's documentation of QC of media is retained
- ___ Adhere to media manufacturer's specifications for intended use
- ___ Document the visual check of all media prior to use (for evidence of contamination, drying, cracking, freezing, etc)
- ___ Read and record temperatures for refrigerator where media is stored and room where DTM cultures are incubated
- ___ Microscope maintenance is performed and recorded

RECORDKEEPING

- ___ Patient test orders include: patient name or identifier; person ordering the test; date of specimen collection; patient age and sex
- ___ Patient test reports include: patient name or identifier; date specimen received; date reported; specimen limitation, if any
- ___ Records are kept for 2 years of lot numbers and expiration dates of media, reagents and dates when placed into use
- ___ The following records are maintained for 2 years: test requests; testing records; reports; quality control and quality assurance activities; proficiency testing or biannual verification of accuracy data.

EXAMPLES OF BIANNUAL VERIFICATION OF ACCURACY ACTIVITIES (2 samples, 2 times/yr)

For microscopic examinations:

- Verify test results by having two analysts review the same specimen and compare findings
- Obtain Kodachrome slides from a reference lab
- Correlate patient test results with clinical presentation

For dermatophyte cultures (growth/no growth only):

- Have reference lab inoculate your media with positive and negative organisms and return to you to incubate and read results.
- Participate in a proficiency testing program
- Obtain stock organisms and inoculate media to test growth/no growth capabilities
- Correlate culture results with KOH prep and clinical findings